WEST Search History

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DATE: Sunday, July 10, 2005

Hide?	Hit Count	
	DB=PGPB; $PLUR=YES$; $OP=OR$	
	L4 (kinase near5 (bead or support)) same peptide	36
	DB=USPT; $PLUR=YES$; $OP=OR$	
	L3 (kinase near5 (bead or support)) same peptide	55
	DB=PGPB,USPT; PLUR=YES; OP=OR	
	L2 (kinase near5 (bead or support)) same peptide	91
	DB=USPT; $PLUR=YES$; $OP=OR$	
	L1 kinase near5 (bead or support)	387

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                 data from INPADOC
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        MAR 02
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        MAR 03
NEWS
      6
        MAR 03
                 MEDLINE file segment of TOXCENTER reloaded
NEWS
      7
NEWS.
     8
        MAR 22
                 KOREAPAT now updated monthly; patent information enhanced
        MAR 22 Original IDE display format returns to REGISTRY/ZREGISTRY
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      10 MAR 22
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                 PATDPASPC - New patent database available
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      11 MAR 22
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      12 APR 04
                 EPFULL enhanced with additional patent information and new
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      14 APR 18
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                 may be affected by a change in filing date for U.S.
                 applications.
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                 U.S. patent records in CA/CAplus
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      20 JUN 06
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                 (Version 8.0 for Windows) now available
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                 and text labels
      25 JUL 01
                 MEDICONF removed from STN
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                 STN Patent Forums to be held in July 2005
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=> kinase (5n) (bead or support)
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=> kinase (5n) (bead or support)
L1 1190 KINASE (5N) (BEAD OR SUPPORT)

=> peptide and 11

L2 122 PEPTIDE AND L1

=> dup rem 12

PROCESSING COMPLETED FOR L2

L3 53 DUP REM L2 (69 DUPLICATES REMOVED)

=> py>2000 and 13

L4 14 PY>2000 AND L3

=> 13 not 14

L5 39 L3 NOT L4

=> t ti 15 1-39

L5 ANSWER 1 OF 39 MEDLINE on STN

TI Serotonin-induced protein kinase C activation in cultured rat heart endothelial cells.

- L5 ANSWER 2 OF 39 MEDLINE on STN
- TI Monoclonal antibodies generated against recombinant ATM support kinase activity.
- L5 ANSWER 3 OF 39 MEDLINE on STN
- TI N-terminal region of P protein of Chandipura virus is responsible for phosphorylation-mediated homodimerization.
- L5 ANSWER 4 OF 39 MEDLINE on STN
- TI Evidence for and against a pivotal role of PI 3-kinase in a neuronal cell survival pathway.
- L5 ANSWER 5 OF 39 MEDLINE on STN
- TI A cell cycle regulated MAP kinase with a possible role in cytokinesis in tobacco cells.
- L5 ANSWER 6 OF 39 MEDLINE on STN
- TI Mechanism of platelet inhibition by nitric oxide: in vivo phosphorylation of thromboxane receptor by cyclic GMP-dependent protein kinase.
- L5 ANSWER 7 OF 39 MEDLINE on STN
- TI Requirement for Rho-mediated myosin light chain phosphorylation in thrombin-stimulated cell rounding and its dissociation from mitogenesis.
- L5 ANSWER 8 OF 39 MEDLINE on STN
- TI Regulation of native Kv1.3 channels by cAMP-dependent protein phosphorylation.
- L5 ANSWER 9 OF 39 MEDLINE on STN
- TI Substrates for protein kinase CK2 in insulin receptor preparations from rat liver membranes: identification of a 210-kDa protein substrate as the dimeric form of endoplasmin.
- L5 ANSWER 10 OF 39 MEDLINE on STN
- TI Effect of calcitonin gene-related **peptide** on sodium absorption through isolated skin of Rana esculenta.
- L5 ANSWER 11 OF 39 MEDLINE on STN
- TI Immunocytochemical localization of protein kinases Yes and Src in amoeboid microglia in culture: association of Yes kinase with vimentin intermediate filaments.
- L5 ANSWER 12 OF 39 MEDLINE on STN
- TI Lipoyl domain-based mechanism for the integrated feedback control of the pyruvate dehydrogenase complex by enhancement of pyruvate dehydrogenase kinase activity.
- L5 ANSWER 13 OF 39 MEDLINE on STN
- TI Differential modulation of bombesin-stimulated phospholipase C beta and mitogen-activated protein kinase activity by [D-Arg1, D-Phe5, D-Trp7,9, Leu11] substance P.
- L5 ANSWER 14 OF 39 MEDLINE on STN
- TI Activation of serine/threonine protein kinases and early growth response 1 gene expression by tumor necrosis factor in human myeloid leukemia cells.
- L5 ANSWER 15 OF 39 MEDLINE on STN
- TI The serum response factor nuclear localization signal: general implications for cyclic AMP-dependent protein kinase activity in control of nuclear translocation.

- L5 ANSWER 16 OF 39 MEDLINE on STN
- TI CD28 signal transduction: tyrosine phosphorylation and receptor association of phosphoinositide-3 kinase correlate with Ca(2+)-independent costimulatory activity.
- L5 ANSWER 17 OF 39 MEDLINE on STN
- TI Casein kinase II mediates multiple phosphorylation of Saccharomyces cerevisiae eIF-2 alpha (encoded by SUI2), which is required for optimal eIF-2 function in S. cerevisiae.
- L5 ANSWER 18 OF 39 MEDLINE on STN
- TI Ro 32-0432, a selective and orally active inhibitor of protein kinase C prevents T-cell activation.
- L5 ANSWER 19 OF 39 MEDLINE on STN
- TI Partial activation of the pyruvate dehydrogenase kinase by the lipoyl domain region of E2 and interchange of the kinase between lipoyl domain regions.
- L5 ANSWER 20 OF 39 MEDLINE on STN
- TI At least two kinases phosphorylate the MPM-2 epitope during Xenopus oocyte maturation.
- L5 ANSWER 21 OF 39 MEDLINE on STN
- TI Insulin receptor serine kinase activation by casein kinase 2 and a membrane tyrosine kinase.
- L5 ANSWER 22 OF 39 MEDLINE on STN
- TI Overexpression of protein kinase C isoenzymes alpha, beta I, gamma, and epsilon in cells overexpressing the insulin receptor. Effects on receptor phosphorylation and signaling.
- L5 ANSWER 23 OF 39 MEDLINE on STN
- TI Mechanistic studies on rhodopsin kinase. Light-dependent phosphorylation of C-terminal peptides of rhodopsin.
- L5 ANSWER 24 OF 39 MEDLINE on STN
- TI Direct photoaffinity-labelling of human deoxycytidine kinase with the feedback inhibitor dCTP.
- L5 ANSWER 25 OF 39 MEDLINE on STN
- TI Electrophoretic purification of the alpha and beta subunits of phosphorylase kinase and evidence in support of the deduced amino acid sequences.
- L5 ANSWER 26 OF 39 MEDLINE on STN
- TI Ultrastructural localization of cyclic adenosine 3',5'-monophosphatedependent protein kinase after adrenocorticotropin stimulation in adrenal cortical tumor cells.
- L5 ANSWER 27 OF 39 MEDLINE on STN
- TI Interleukin 2 and diacylglycerol stimulate phosphorylation of 40 S ribosomal S6 protein. Correlation with increased protein synthesis and S6 kinase activation.
- L5 ANSWER 28 OF 39 MEDLINE on STN
- TI Altered phosphoglycerate kinase from old rat muscle shows no change in primary structure.
- L5 ANSWER 29 OF 39 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI Poly(ADP-ribose) modulates the properties of MARCKS proteins.

- L5 ANSWER 30 OF 39 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Affinity purification of recombinant proteins fused to calmodulin or to calmodulin-binding peptides
- L5 ANSWER 31 OF 39 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Application of the one-bead one-compound combinatorial library method in protein tyrosine kinase and cell surface receptor research
- L5 ANSWER 32 OF 39 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Identification and characterization of a novel **peptide** substrate for P60c-src protein tyrosine **kinase** using a one-**bead** one-**peptide** combinatorial **peptide** library method
- L5 ANSWER 33 OF 39 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Template directed cyclization of support-bound peptides.
- L5 ANSWER 34 OF 39 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Activation of serine/threonine protein kinases and early growth response 1 gene expression by tumor necrosis factor in human myeloid leukemia cells
- L5 ANSWER 35 OF 39 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Method for the detection of phosphotyrosine residues
- L5 ANSWER 36 OF 39 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Analysis and mapping of plastin phosphorylation.
- L5 ANSWER 37 OF 39 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Regulation of native Kv1.3 channels by cAMP-dependent protein phosphorylation.
- L5 ANSWER 38 OF 39 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Synergistic activation of a G protein-coupled receptor kinase by G protein $\beta\gamma$ subunits and mastoparan or related peptides.
- L5 ANSWER 39 OF 39 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Inhibition of neutrophil superoxide formation by 1-(5-isoquinolinesulfonyl)-2-methylpiperazine (H-7), an inhibitor of protein kinase-C.

=> 15 and bead

L6 4 L5 AND BEAD

=> d ibib abs 16 1-4

L6 ANSWER 1 OF 4 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN

ACCESSION NUMBER: 1998:342207 BIOSIS DOCUMENT NUMBER: PREV199800342207

TITLE: Poly(ADP-ribose) modulates the properties of MARCKS

proteins.

AUTHOR(S): Schmitz, Arndt A. P.; Pleschke, Jutta M.; Kleczkowska,

Hanna E.; Althaus, Felix R. [Reprint author]; Vergeres, Guy

[Reprint author]

CORPORATE SOURCE: Dep., Biophysical Chem., Biozentrum, Univ. Basel,

Klingelbergstrasse 70, CH-4056 Basel, Switzerland Biochemistry (June 30, 1998) Vol. 37, No. 26, pp.

SOURCE: Biochemistry, (June 30, 1998) Vol. 37, No. 26, pp.

9520-9527. print.

CODEN: BICHAW. ISSN: 0006-2960.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 13 Aug 1998

Last Updated on STN: 10 Sep 1998

In mammalian cells, the formation of DNA strand breaks is accompanied by AB synthesis of poly(ADP-ribose). This nucleic acid-like homopolymer may modulate protein functions by covalent and/or noncovalent interactions. Here we show that poly(ADP-ribose) binds strongly to the proteins of the myristoylated alanine-rich C kinase substrate (MARCKS) family, MARCKS and MARCKS-related protein (also MacMARCKS or F52). MARCKS proteins are myristoylated proteins associated with membranes and the actin cytoskeleton. As targets for both protein kinase C (PKC) and calmodulin (CaM), MARCKS proteins are thought to mediate cross-talk between these two signal transduction pathways. Dot blot assays show that poly(ADP-ribose) binds to MARCKS proteins at the highly basic effector domain. Complex formation between MARCKS-related protein and CaM as well as phosphorylation of MARCKS-related protein by the catalytic subunit of PKC are strongly inhibited by equimolar amounts of poly(ADP-ribose), suggesting a high affinity of poly(ADP-ribose) for MARCKS-related protein. Binding of MARCKS-related protein to membranes is also inhibited by poly(ADP-ribose). Finally, poly(ADP-ribose) efficiently reverses the actin-filament bundling activity of a peptide corresponding to the effector domain and inhibits the formation of actin filaments in vitro. Our results suggest that MARCKS proteins and actin could be targets of the poly(ADP-ribose) DNA damage signal pathway.

L6 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1998:440824 CAPLUS

DOCUMENT NUMBER: 129:211222

TITLE: Application of the one-bead one-compound

combinatorial library method in protein tyrosine

kinase and cell surface receptor research

AUTHOR(S): Lam, K. S.; Lou, Q.; Wu, J.; Leftwich, M.; Mckay, R.

T.; Rychetsky, L.; Phan, H.; Joe, J.; Chen, M. -L.;

Liu-Stevens, R.; Zhao, Y.; Salmon, S. E.

CORPORATE SOURCE: Arizona Cancer Center, Department of Medicine,

University of Arizona, Tucson, AZ, 85724, USA

SOURCE: Peptides: Biology and Chemistry, Proceedings of the

Chinese Peptide Symposium, 4th, Chengdu, Peop. Rep. China, July 21-25, 1996 (1998), Meeting Date 1996, 55-58. Editor(s): Xu, Xiao-Jie; Ye, Yun-Hua; Tam,

James P. Kluwer: Dordrecht, Neth.

CODEN: 66KJAP

DOCUMENT TYPE: Conference LANGUAGE: English

The "one-bead one-compound" combinatorial library method is extremely versatile and can be used to discover ligands for various mol. targets. Assays can be developed such that a specific biol. or phys. property can be detected. These assays, whether on-bead or in solution phase can easily be adapted to the "one-bead one-compound" library concept in e.g. protein tyrosine kinase and cell surface receptor research. Thus far, this specific combinatorial library method has proven

REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1996:695882 CAPLUS

DOCUMENT NUMBER: 126:3618

TITLE: Identification and characterization of a novel

to be very useful in both basic research and drug discovery.

peptide substrate for P60c-src protein

tyrosine kinase using a one-bead one-peptide combinatorial peptide

library method

AUTHOR(S): CORPORATE SOURCE: Lam, K. S.; Lou, Q.; Wu, J.; Salmon, S. E.; Phan, H. Arizona Cancer Center, University Arizona, Tucson, AZ,

85724, USA

SOURCE:

Peptides: Chemistry, Structure and Biology,

Proceedings of the American Peptide Symposium, 14th, Columbus, Ohio, June 18-23, 1995 (1996), Meeting Date 1995, 287-289. Editor(s): Kaumaya, Pravin T. P.; Hodges, Robert S. Mayflower Scientific: Kingswinford,

UK.

CODEN: 63NTAF

DOCUMENT TYPE:

Conference

LANGUAGE:

English

We have successfully applied a one-bead one-peptide combinatorial peptide library method for identification of linear peptide substrate motifs for cAMP-dependent protein kinase (a serine/threonine protein kinase) and for P60c-src protein tyrosine kinase (PTK). In this method, we first incubated the peptide-bead library with $[\gamma-32P]$ ATP and the protein kinase. After incubation, the beads were washed thoroughly with high salt buffer followed by heating with 1.0 M HCl for 5 min to remove all the non-covalent $[\gamma-32P]$ ATP binding and washed thoroughly again. The beads were then suspended in molten 1.5% (w/v) agarose and plated on a glass plate. The bead-containing gel was then air-dried to form a film and exposed to an X-ray film. Autoradiog. was then used to localize the [32P]-labeled beads. The beads corresponding to the autoradiog. spots were removed and suspended in molten agarose solution again for secondary plating. With this dilution, single [32P]-labeled beads could be isolated

ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1993:209001 CAPLUS

DOCUMENT NUMBER:

118:209001

TITLE:

Method for the detection of phosphotyrosine residues

INVENTOR(S):

Ziltener, Hermann J.

PATENT ASSIGNEE(S):

Can.

for microsequencing.

SOURCE:

PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9303377	A1	19930218	WO 1992-CA328	19920730

RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, MC, NL, SE PRIORITY APPLN. INFO.: US 1991-739141 A 19910731

A sensitive and rapid method for detecting phosphotyrosine residues uses antiphosphotyrosine antibody in a particle concentration fluorescence immunoassay. This immunoassay can be used to measure the activity of and screen for a protein tyrosine kinase, a protein tyrosine phosphatase, and their modulators and substrates. Fluoricon 0.8-µm diameter carboxyl-activated polystyrene particles were coupled with myelin basic protein or a peptide derived from protein tyrosine kinase p56lck. Protein tyrosine kinase p56lck was assayed by adding a mixture of substrate-coated particles in Tris-HCl buffer containing ATP and MnCl2 to wells of a filtration plate, adding sample to the wells, incubating at 37° for 15 min, draining the wells, washing with buffer to remove

kinase, adding anti-phosphotyrosine monoclonal antibody, and detecting bound antibody by particle concentration fluorescence immunoassay.

=> d his

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-2.19

-2.19

JUL 2005					
1190 KINASE (5N) (BEAD OR SUPPORT)					
L2 122 PEPTIDE AND L1					
L3 53 DUP REM L2 (69 DUPLICATES REMOV	3 DUP REM L2 (69 DUPLICATES REMOVED)				
4 14 PY>2000 AND L3					
L5 39 L3 NOT L4					
L6 4 L5 AND BEAD					
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION			

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